## IN THE CLAIMS:

Please amend claims 41, 42 and 46, without prejudice, as follows:

Claims 1-40 (cancelled).

Claim 41 (currently amended): A method of <u>treating</u> inhibiting-HIV replication infection in a mammal, <u>wherein said HIV infection is modulated by a CCR5 receptor in said mammal</u>, comprising administering to said mammal an effective amount of a compound of formula I or a pharmaceutically acceptable salt or solvate thereof:

wherein  $R^1$  is  $C_{3\cdot6}$  cycloalkyl optionally substituted by one or more fluorine atoms, or  $C_{1\cdot6}$  alkyl optionally substituted by one or more fluorine atoms, or  $C_{3\cdot6}$  cycloalkylmethyl optionally ring-substituted by one or more fluorine atoms; and

R<sup>2</sup> is phenyl optionally substituted by one or more fluorine atoms.

Claim 42 (currently amended): A method of <u>treating</u> inhibiting-HIV replication-infection in a mammal, <u>wherein said HIV infection is modulated by a CCR5 receptor in said mammal</u>, comprising administering to said mammal an effective amount of a compound selected from the group consisting of:

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}cyclobutanecarboxamide;

*N*-{(1*S*)-3-[3-(3-Isopropyl-5-methyl-4*H*-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}cyclopentanecarboxamide;

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}-4,4,4-trifluorobutanamide;

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}-4,4-difluorocyclohexanecarboxamide; and

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-(3-fluorophenyl)propyl}-4,4-difluorocyclohexanecarboxamide;

or a pharmaceutically acceptable salt or solvate thereof.

Claims 43-45 (Cancelled).

Claim 46 (currently amended): A method of <u>treating inhibiting-HIV replication-infection</u> in a mammal, <u>wherein said HIV infection is modulated by a CCR5 receptor in said mammal</u>, comprising administering to said mammal an effective amount of N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}-4,4-difluorocyclohexanecarboxamide or a pharmaceutically acceptable salt or solvate thereof.